AMENDMENTS TO THE SPECIFICATION

Page 2, paragraph at lines 6-11:

said method comprising adding to an imidazolidinethione having formula

one of: (i) CHR5-CHR6-C(Y)ZR7 CHR5-CR6-C(Y)ZR7; and (ii) R8N-C=W to form a reaction mixture; wherein the reaction mixture is substantially free of solvent.

Page 4, paragraph at lines 6.8:

(b) adding to the imidazolidinethione, without isolation of the imidazolidinethione, one of (i) CHR5-CHR5-C(Y)ZR7 CHR5-CR6-C(Y)ZR7; (ii) $R^{10}R^{11}C=0$ and $R^{12}NH_2$; (iii) $R^{10}R^{11}C=NR^{12}$; and (iv) $R^8N=C=W$.

Page 7, paragraph at lines 5-20:

In one embodiment of the invention, the imidazolidinethione reacts with CHR6-CHR6-C(Y)ZR7 CHR5-CR6-C(Y)ZR7 or R8N=C=W substantially in the absence of a solvent. A solvent is any liquid other than the reactants or products of this reaction. Preferably, the reaction mixture contains no more than 5% of solvent by weight, more preferably no more than 2%, more preferably the reaction mixture contains no solvent. Elimination of the solvent increases the efficiency of the process by reducing the cost and the reaction volume. Preferably, the reaction with CHR5=CHR6-C(Y)ZR7 CHR5=CR6-C(Y)ZR7 or R8N=C=W is performed at a temperature from 50°C to 180°C, more preferably from 60°C to 170°C, and most preferably from 90°C to 130°C. The reaction may be followed by well-known methods to determine reaction completion, e.g., IR spectroscopy. Typically, the

reaction is complete in 0.5 to 4 hours. Substitution of acrylate occurs on the thioamide nitrogen or sulfur atom, thereby producing a -CHR⁵-CHR⁶-C(Y)ZR⁷ group as B¹ or B², respectively. In contrast, substitution of R⁸N=C=W occurs on the amine nitrogen atom of the imidazolidinethione ring, thereby producing a -C(W)NHR⁸ group as B³.

Page 8, paragraph at lines 8-18:

In one embodiment of the invention, an imidazolidinethione is prepared, resulting in a reaction mixture containing the imidazolidinethione, a solvent (typically water or a partially aqueous solvent), and possibly starting materials and byproducts. In this embodiment, one of: (i) CHR5=CHR6-C(Y)ZR7 CHR5=CR6-C(Y)ZR7 CHR5=CR6-C(Y)ZR7; (ii) R¹0R¹¹C=O and R¹²NH2; (iii) R¹0R¹¹C=NR¹²; and (ii) R³N=C=W is added to the reaction mixture without isolation of the imidazolidinethione. Addition of one of these reagents directly to the imidazolidinethione reaction mixture increases the efficiency of the process by eliminating a costly purification step. In one preferred embodiment, the water is partially or substantially completely removed from the reaction mixture prior to addition of one of the aforementioned reagents.